



Cure SMA Walk-n-Roll Registration

Sunday, October 3, 2010, 12:30 pm, Winchester, Massachusetts

Name(s): _____

Walking in honor/memory of: _____

We: ☐ cannot join you but have enclosed a donation. (Or you may donate online.)

☐ would like to register ____ participants

of T-shirts: ____Adult XL ____Adult L ____ Adult M ____Adult S ____Youth XL ____Youth L

Address: _____ Phone: _____

City/State/Zip _____ E-mail: _____

We would like to receive email updates from the New England Chapter of FSMA: ____Yes ____ No

You may bring this form with you to the walk or mail it to BJ Mirabile,

43 Jefferson Rd., Winchester, MA, 01890; or you may donate online or register and collect pledges online at <http://www.fsma.org/lynchwalkforacure>.

•Please make checks payable to:

Families of SMA.

•All donations are fully tax deductible.

•The registration fee is \$20 per person
Pledges collected can be applied to registration.

•You can donate online, or register and collect online pledges at
www.fsma.org/lynchwalkforacure

•For more information contact BJ at
781-721-7862 or bjmirabile@verizon.net

Sponsor Name	Address or Phone	Donation
Thank you for your contribution.	Total:	

Participants please sign: I hereby acknowledge that I am physically able to undertake Walk-n-Roll and waive any and all claims arising out of Walk-n-Roll which I might assert against Families of SMA, their directors and volunteers.

Participant's Signature

**Lynch Elementary School
10 Brantwood Road
Winchester, MA**

Spinal Muscular Atrophy...

...is an inherited disease which destroys the nerves that command voluntary muscle movement.

...affects a child's ability to crawl, walk, control the head, and even swallow and breathe.

...is carried by one in 40 people and afflicts one in 6,000 babies born.

...is the number one genetic killer of children under the age of two.

... currently has no treatment or cure.

Families of SMA...

...is the largest non-profit organization dedicated solely to eradicating SMA.

...helps support families coping with SMA and educates the public and professional community about SMA.

...promotes and funds basic research, drug discovery and development, and clinical trials.

Reasons for Hope...

FSMA was awarded a multi-million dollar grant from the NINDS to continue development of the Tetracycline drug candidate with Paratek Pharmaceuticals and...

...FSMA developed drug candidate Quinazoline495, which received Orphan Drug Designation from the FDA, and was licensed to Repligen Corporation for clinical development as a potential treatment for SMA.

Kelvyn A.
Tyler A.
Arcand A.
Andy A.
Jack B.
Anwar B.
Peter B.
Alex B.
Caitlyn B.
Julia B.
Aleksandar B.
Abby B.
John B.
Nolan B.
Rachel B.
Angelo C.
Diana C.
Dillan C.
Jasmine C.
Olivia C.
Rico C.
Edward C.
Tommy D.
Jackson D.
Grace D.
Elizabeth E.
Spencer E.
John E.
Rhiannon F.
Carlina F-M.
Maggie F.
Skyler G.
Anthony G.
Allison G.
Cristian H.
Tucker H.
Berke H.
Justice H.
VanTung H.

Mae Z.

Jeremy H.
Justin J.
Josh J.
Jason K.
Justin K.
Tyler L.
LingLong L.
Jonathan L.
Benny L.
Will M.
Jonathan M.
Christy M.
Lauren M.
Andy M.
Charley M.
Katie M.
Stefano M.
Shameem M.
Meghan M.
Steven O.
Bryan P.
Jeremy P.
Katie P.
Jason P.
Alex P.
Chloe R.
Kate R.
Matt S.
Allen S.
Valeriy S.
Dashannah S.
Aiden S.
Adrian S.
Andrew T.
Eric W.
Jerry W.
Michael X.
Alfie Y.
Ben Z.

Lynch School 5th graders Walking for a Cure for SMA to fund research for Spinal Muscular Atrophy



Sunday, October 3, 2010
rain or shine
Lynch Elementary School
Winchester, MA

registration begins 12:30 pm
walk starts 1:30 pm

Join us for family fun, refreshments,
and great raffle items.
~2 miles walk.

